## AMENDMENT TO THE SPECIFICATION

Kindly amend the specification at page 11, line 7, through page 12, line 9, as follows:

Pluripotent mesenchymal cells derived from UCB can be administered directly and induced to differentiate by contact with tissue in vivo or induced to differentiate into a desired cell type, e.g., mesenchymal cells, hematopoietic cells, neural cells, or endothelial cells, etc., using in vitro or ex vivo methods before their administration. Such predisposition of progeny of pluripotent mesenchymal cells derived from UCB has the potential to shorten the time required for complete differentiation once the cells have been administered to the patient. Techniques for the differentiation of pluripotent cells into cells of a particular phenotype are known in the art, such as those described in U.S. Patent Nos. 5,486,359; 5,591,625; 5,736,396; 5,811,094; 5,827,740; 5,837,539; 5,908,782; 5,908,784; 5,942,225; 5,965,436; 6,010,696; 6,022,540; 6,087,113; 5,858,390; 5,804,446; 5,846,796; 5,654,186; 6,054,121; 5,827,735; and 5,906,934, which describe the transformation of pluripotent cells. For example, Rodgers et al. (U.S. Patent. No. 6,335,195), describes methods for the ex vivo culturing of hematopoietic and mesenchymal pluripotent cells and the induction of lineage-specific cell proliferation and differentiation by growth in the presence of angiotensinogen, angiotensin I (AI), AI analogues, AI fragments and analogues thereof, angiotensin II (AII), AII analogues, AII fragments or analogues thereof, or AII AT<sub>2</sub>-type 2 receptor agonists, either alone or in combination with other growth factors and cytokines. In an embodiment, the pluripotent cells of the invention can be induced in vitro to differentiate into pancreatic cells, and in particular pancreatic islet cells, by using, e.g., techniques known in the art (see, e.g., Yang et al., Proc. Nat. Acad. Sci. USA 99: 8078-83, 2002; Zulewski et al., Diabetes 50: 521-33, 2001; and Bonner-Weir et al., Proc. Nat. Acad. Sci. USA 97: 7999-8004, 20002001). Art-known techniques can also be used to induce the pluripotent cells of the invention to differentiate in vitro into hepatic cells (see, e.g., Lee et al., Hepatology 40: 1275-1284, 2004), neuronal cells (see, e.g., Thondreau et al., Differentiation 319-322-326, 2004), or endothelial cells (see, e.g., Kassem et al., Basic Clin. Pharmacol. & Toxicol. 95:209-214, 2004; and Pittenger and Martin, Circ. Res. 95:9-20, 2004). Optionally, a differentiating

agent may be co-administered or subsequently administered to the subject to promote stem cell differentiation *in vivo*.